BOSTER BIOLOGICAL TECHNOLOGY
3942 B Valley Ave, Pleasanton, CA 94566
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Anti-Zebrafish NME7 Antibody Picoband®

Catalog Number: AZA0A8N7UZE3

About NME7

Predicted to enable 3'-5' exonuclease activity; ATP binding activity; and nucleoside diphosphate kinase activity. Predicted to be involved in CTP biosynthetic process; GTP biosynthetic process; and UTP biosynthetic process. Predicted to be located in nucleus and spindle. Predicted to be active in axonemal microtubule and centrosome. Is expressed in several structures, including digestive system; eye; female organism; gonad; and male organism. Orthologous to human NME7 (NME/NM23 family member 7).

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Overview

Product Name	Anti-Zebrafish NME7 Antibody Picoband®
Reactive Species	Zebrafish
Description	Boster Bio Anti-Zebrafish NME7 Antibody Picoband® catalog # AZA0A8N7UZE3. Tested in WB, IHC applications. This antibody reacts with Zebrafish. The brand Picoband indicates this is a premium antibody that guarantees superior quality, high affinity, and strong signals with minimal background in Western blot applications. Only our best-performing antibodies are designated as Picoband, ensuring unmatched performance.
Application	IHC, WB
Clonality	Polyclonal
Formulation	Each vial contains 4 mg Trehalose, 0.9 mg NaCl, 0.2 mg Na2HPO4.
Storage Instructions	At -20°C for one year from date of receipt. After reconstitution, at 4°C for one month. It can also be aliquotted and stored frozen at -20°C for six months. Avoid repeated freezing and thawing.
Host	Rabbit
Uniprot ID	A0A8N7UZE3

Technical Details

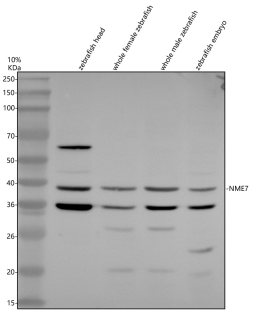
Immunogen	E.coli-derived Zebrafish NME7 recombinant protein (Position: K29-E215).
Form	Lyophilized

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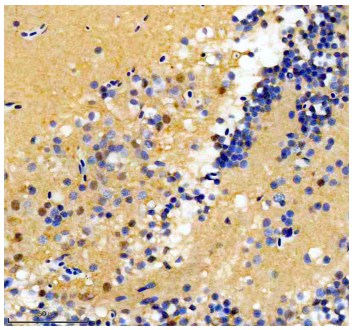
Concentration	Adding 0.2 ml of distilled water will yield a concentration of 500 ug/ml.
Purification	Immunogen affinity purified.
Suggested Dilutions	Western blot, 0.25-0.5 ug/ml, Zebrafish Immunohistochemistry (Paraffin-embedded Section), 2-5 ug/ml, Zebrafish

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Anti-Zebrafish NME7 Antibody Picoband® (AZA0A8N7UZE3) Images



Western blot analysis of NME7 using anti-NME7 antibody (AZA0A8N7UZE3). Electrophoresis was performed on a 10% SDS-PAGE gel at 80V (Stacking gel) / 120V (Resolving gel) for 2 hours. The sample well of each lane was loaded with 30 ug of sample under reducing conditions. Lane 1: zebrafish head tissue lysates, Lane 2: whole female zebrafish lysates, Lane 3: whole male zebrafish lysates, Lane 4: zebrafish embryo lysates.After electrophoresis, proteins were transferred to a nitrocellulose membrane at 150 mA for 50-90 minutes. Blocked the membrane with 5% non-fat milk/TBS for 1.5 hour at RT. The membrane was incubated with rabbit anti-NME7 antigen affinity purified polyclonal antibody (AZA0A8N7UZE3) at 0.5 ug/mL overnight at 4°C, then washed with TBS-0.1%Tween 3 times with 5 minutes each and probed with a goat anti-rabbit IgG-HRP secondary antibody at a dilution of 1:5000 for 1.5 hour at RT. The signal is developed using an ECL Plus Western Blotting Substrate (Catalog # AR1196-200) with Tanon 5200 system. A specific band was detected for NME7 at approximately 39 kDa. The expected band size for NME7 is at 42 kDa.



IHC analysis of NME7 using anti-NME7 antibody (AZA0A8N7UZE3). NME7 was detected in a paraffin-embedded section of zebrafish brain tissue. Heat mediated antigen retrieval was performed in EDTA buffer (pH 8.0, epitope retrieval solution). The tissue section was blocked with 10% goat serum. The tissue section was then incubated with 2 ug/ml rabbit anti-NME7 Antibody (AZA0A8N7UZE3) overnight at 4°C. Peroxidase Conjugated Goat Anti-rabbit IgG was used as secondary antibody and incubated for 30 minutes at 37°C. The tissue section was developed using HRP Conjugated Rabbit IgG Super Vision Assay Kit (Catalog # SV0002) with DAB as the chromogen.

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